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Electronic and chiroptical properties of chiral cycloiridiated complexes bearing helicenic NHC ligands

Nora Hellou,^a Claire Jahier-Diallo,^{b,f} Olivier Baslé,^b Monika Srebro-Hooper,^c Loïc Toupet,^a Thierry Roisnel,^a Elsa Caytan,^a Christian Roussel,^d Nicolas Vanthuyne,^d Jochen Autschbach,^{*e} Marc Mauduit,^b and Jeanne Crassous^{*a}

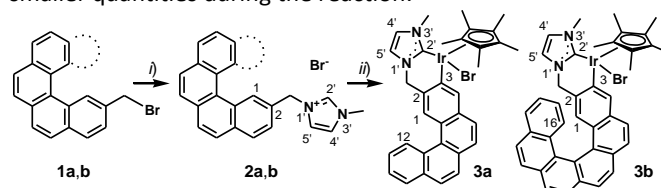
The first helicene-NHC-iridium complexes have been prepared in enantiopure forms. Their stereochemical, electronic, and chiroptical features were characterized experimentally and via first-principles calculations, highlighting the helical and iridium-centered chirality.

N-heterocyclic carbene (NHC) transition-metal (TM) complexes have undergone rapid development in the past two decades due to their important applications in catalysis, medicinal chemistry, and molecular materials.^{1,2} Notably, the high structural modularity of NHCs has enabled the design of a large variety of chiral architectures, which gave rise to highly stereoselective organometallic catalysts.³ On the other hand, chiral NHC-TM systems may combine optical activity with other desired properties such as emission, electroactivity and high stability, which make them attractive candidates as multifunctional optoelectronic materials.⁴ Nevertheless, studies exploring the chiroptical properties of NHC-based TM-complexes are still very rare.⁵ Furthermore, while the coordination of helicenic ligands to TM has recently led to organometallic species with unique physico-chemical properties,^{6,7} this approach has never been considered in NHC chemistry.⁸

This communication describes the preparation of the first cycloiridiated NHC complexes bearing either an achiral [4]helicene (*i.e.* a benzophenanthrene) or a chiral [6]helicene moiety. In complexes **3a,b**, the Ir(III) is a stereogenic center as well.⁹ The stereochemical features, along with electronic and chiroptical properties, *i.e.* electronic circular dichroism (ECD) and molar rotation (MR), are examined both experimentally and theoretically.

The two targeted systems are cycloiridiated complexes (**3a,b**) of general formula (**4a,b**)IrBrCp* where Cp* = pentamethylcyclopentadienyl, and **4a,b** are the LX-type NHC ligands derived from either a [4]helicene- (**2a**) or a [6]helicene-based unit (**2b**) grafted at position 2 by a 1-methylene-3-

methylimidazolium bromide (Scheme 1). Imidazolium salt **2a** was prepared from 2-bromomethylene[4]helicene^{10a} **1a** by means of nucleophilic substitution with 3-methylimidazolium. The subsequent cyclometallation reaction in refluxing acetonitrile, using [Cp*IrCl₂]₂ as the iridium source and a base, such as for instance NaOAc, yielded complex **3a** with 57% yield after crystallization. The crystallization step was necessary to remove the chlorido complex, *i.e.* (**4a**)IrClCp* that is formed in smaller quantities during the reaction.^{11,*}



Scheme 1. Synthesis of "chiral at metal" Ir(III) complexes **3a,b**. *i*) Methylimidazole, acetone, reflux, 15 hrs, 66-71%; *ii*) [Cp*IrCl₂]₂, NaOAc, CH₃CN, 80°C, 15 hrs, 35-57% after crystallization.

Compound **3a** was fully characterized by NMR spectroscopy, mass spectrometry and X-ray crystallography (see Electronic Supporting Information, ESI). For instance, its ¹H NMR spectrum displays the typical deshielded H¹ and H¹² helicenic protons at 8.79 and 9.20 ppm respectively, while the H^{2'}-imidazolium and H³-helicene signals have disappeared. The bridging methylene group appears as an AB system around 5 ppm (with a 14 Hz coupling constant) similarly to analogous cycloiridiated complexes.¹¹ Furthermore, two types of methyl groups are observed at 3.97 and 1.76 ppm for the 3'-methylimidazole and the Cp*, respectively.

Racemic complex **3a** crystallized in the centro-symmetric P₂₁/n space group. X-ray crystallographic studies (Figure 1) revealed that the cyclometallation occurred preferentially at position 3 rather than at position 1, certainly due to less steric congestion. The Ir-C² distance of 1.998 Å is characteristic of an Ir-NHC bond while the Ir-C³ bond-length of 2.047 Å ascertains that the cyclometallation occurred.¹¹ Two stereogenic elements are found in the solid state, *i.e.* the Ir center and the [4]helicene unit, with only two enantiomeric pairs, *i.e.* (*M*,*R*_{Ir}) and (*P*,*S*_{Ir}). The other diastereomers are not present. The [4]helicene moiety displays a helical angle of 30°. Note, however, that in solution only the stereogenic Ir center is stable since the [4]helicene fragment readily epimerizes (*vide infra*). The 6-membered iridacycle adopts a boat conformation and π-π stacking interactions are observed between the Cp* and the [4]helicene of two homochiral pairs with the same (*P*,*S*_{Ir}) or (*M*,*R*_{Ir}) configurations (see Figure 1).

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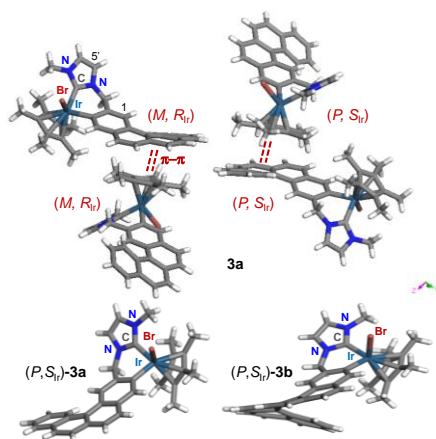


Figure 1. X-ray structure of racemic (*M,R_{ir}*)/(*P,S_{ir}*)-**3a** and its supramolecular organization. X-ray structure of enantiopure (*P,S_{ir}*)-**3a** and (*P,S_{ir}*)-**3b**.

The conformational equilibrium of the 6-membered iridacycle was investigated by density functional theory (DFT) calculations (BP functional with a continuum solvent model for CH₂Cl₂). For the (*P,S_{ir}*)-**3a** diastereomer, two stable boat-like conformations were found and optimized. However, the conformer with the CH₂ group pointing towards Cp* (as in the XRD structure of Figure 1) is significantly lower in energy (by 13.1 kcal/mol) than the one with CH₂ pointing towards the Br ligand (see ESI), and therefore latter can be neglected. (*P,R_{ir}*)-**3a** affords the same strong conformational preference (14.3 kcal/mol). Furthermore, NOESY experiments showed clear correlation peaks between one methylenic proton and H¹ / H⁵ from helicene / NHC units, respectively, while the other CH₂ proton correlates with the Cp* (see ESI), thus confirming the blocked conformation of the iridacycle.

The enantiomeric resolution of this new type of neutral helicenic "chiral at metal"^{6c,9,12} Ir(III) complex was achieved by HPLC over a chiral stationary phase (see ESI). The *R_{ir}* and *S_{ir}* enantiomers of **3a** were obtained with good enantiomeric excesses (*ee*'s of 93 and 96%, respectively) after separation over the Chiralpak® IC column (using heptane/ethanol/chloroform mixture as the eluent at room temperature, r.t.). Kinetic studies, based on the evolution of the ECD at 254 nm with time at 60°C (see ESI, Figure S2, page S7), estimate the racemization barrier of (-)-**3a** around 110 kJ/mol and a half-life time of ca. 2 hours at 62°C in chloroform. According to the DFT calculations, (*P,S_{ir}*)- and (*P,R_{ir}*)-**3a** have a negligible difference in energy (with a very slight preference for (*P,S_{ir}*), see ESI), indicating that **3a** may exist as an equal mixture of both diastereomers. To avoid potential decomposition and partial racemization, the complexes were stored at low temperature and in the dark.

The chiroptical properties (electronic circular dichroism (ECD) and molar rotation (MR))⁵ of *R_{ir}* and *S_{ir}*-**3a** were then studied. The two mirror-image ECD spectra are depicted in Figure 2. For example, the *S_{ir}*-**3a** enantiomer displays a positive ECD-active band at 280 nm ($\Delta\epsilon = +36$), two negative ones at 306 ($\Delta\epsilon = -18$) and 360 nm ($\Delta\epsilon = -10$) and a very weak positive one ($\Delta\epsilon = +7$) at 413 nm. Overall the ECD of "chiral at Ir" complex **3a** is relatively weak. The MR is also low (*S_{ir}*(+)-**3a**:

$[\phi]_D^{23} = +230 (\pm 7\%)$, $C 1.8 \cdot 10^{-3}$ M, CH₂Cl₂). The *S_{ir}*(+) and *R_{ir}*(-) absolute configurations were determined by X-ray crystallography of enantiopure (+)-**3a** which crystallized in the P2₁2₁2₁ space group (Figure 1). Time-dependent DFT (TDDFT, BHLYP / PBE0 functional for MR / ECD, CH₂Cl₂ solvent model) calculations of the chiroptical properties were also performed.¹³ The Boltzmann-average of the calculated (*P,S_{ir}*)- and (*M,S_{ir}*)-**3a** MRs is 284 and agrees very well with the experiment.

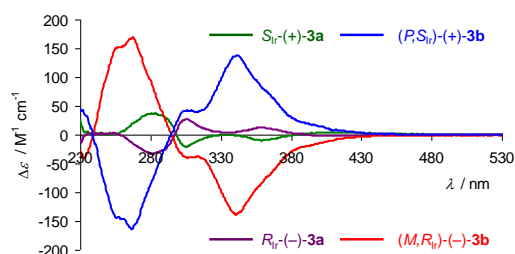


Figure 2. Electronic circular dichroism spectra of complexes *R_{ir}*- and *S_{ir}*-**3a** and (*P,S_{ir}*)- and (*M,R_{ir}*)-**3b** in CH₂Cl₂ at 20°C. ($C \sim 5 \cdot 10^{-5} - 2 \cdot 10^{-4}$ M).

The sign pattern of the experimentally observed ECD bands, and their relative intensities, are also correctly reproduced in the simulated ECD spectrum (see the ESI for the full set of calculated data and spectral analyses). For instance, in the computed spectrum of *S_{ir}*-**3a**, for increasing energy / decreasing wavelength a very weak positive band is followed by two negative bands and a positive one (Figure 3). It should be stressed that the simulated ECD spectrum is obtained from an average of optimized structures for the *P* and *M* helicities of the ligand, with concomitant cancellation of nearly opposite rotatory strengths. Therefore, the averages are sensitive to the energy difference between the (*P,S_{ir}*)- and (*M,S_{ir}*)-**3a** stereoisomers, the functional used in the TDDFT computations, and other theoretical details (see ESI). Similar considerations apply to the calculated MR. However, the lowest-energy electronic excitation always appears with positive rotatory strength and is indicative of the configuration around the metal (*vide infra*). The lowest-energy part of *S_{ir}*-**3a** spectrum results from two excitations (calculated ca. 410 / 380 nm for no. 1 / 2), with low-magnitude rotatory strengths of opposite sign (+ / - for no. 1 / 2). According to a molecular orbital (MO) decomposition of the spectra, the dominant contributions to these excitations arise from HOMO – LUMO+2 and HOMO-1 – LUMO+2 orbital pairs. These orbitals are centered at and around the metal, which rationalizes why the excitations are influenced by the absolute configuration around Ir. The excitations represent a mixture of ligand-field (LF) as well as ligand-to-metal (LM) and ML charge-transfer (CT) character. Overall, the theoretical analysis highlights the role of the metal center and the carbene-helicene moiety in determining optical activity of these novel compounds.

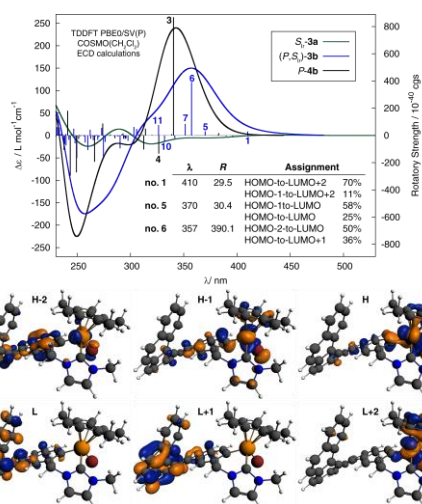


Figure 3. Top: Comparison of the simulated ECD spectra of the Ir-based NHC-helicene complex S_{ir} -**3a**, of the complex (P,S_{ir}) -**3b** and its parent NHC-helicene ligand P -**4b**. No spectral shift has been applied. Calculated excitation energies and rotatory strengths indicated as 'stick' spectra. Numbered excitations correspond to those analyzed in detail. Inset: Selected dominant excitations and occupied-to-unoccupied MO pair contributions of the (P,S_{ir}) -**3b**. Wavelength λ in nm, rotatory strength R in 10^{-40} cgs. Bottom: Selected MOs involved in the ECD-active bands of **3b**. See also ESI.

In order to obtain a configurationally stable helicene derivative, the [6]helicene-based methylimidazolium bromide **2b** was prepared from 2-bromomethylene[6]helicene^{10b} **1b** (Scheme 1). The Ir(III) complex **3b** was synthesized from **2b** under similar conditions as **3a**, and obtained in 35% isolated yield. Here again, a crystallization step was required to remove the chlorido complex (**4b**)IrClCp*.[†] Cycloiridiated complex **3b** was fully characterized and ¹H NMR analyses showed the same general characteristics as **3a** (ESI) with a typical shielded H¹ and H¹⁶ protons of the [6]helicene core (at 7.15 and 7.84 ppm, respectively). In addition, the NMR evidenced only one diastereoisomer which was further identified by X-ray studies as $(P,S_{ir})/(M,R_{ir})$ -**3b**. Furthermore, upon crystallization from a CH₂Cl₂/pentane mixture, racemic complex **3b** spontaneously resolved into homochiral crystals ($P2_12_12_1$ space group). The cyclometallation occurred again at the less hindered position 3 and the Ir-C^{2'} (2.016 Å) and Ir-C³ (2.064 Å) bond-lengths are characteristic of an Ir-NHC cyclometallated complex.¹¹ The [6]helicene unit displays a helical angle of 50°, a typical value for configurationally stable [6]helicene derivatives.^{6,7} Similarly to **3a**, the 6-membered iridacycle adopts a stable boat-like conformation with different correlation peaks for the two CH₂ protons in the ROESY spectrum (see ESI), confirming the presence of one blocked conformation of the iridacycle. Calculations corroborate this finding with an energetic preference of the conformation with CH₂ group pointing towards Cp* by ca. 14 kcal/mol (see ESI).

Since the crystals were too small to achieve Pasteur's resolution by picking enantiopure crystals manually, the chiral HPLC resolution was carried out. Enantioenriched samples (*ee*'s ~ 92-97%) were obtained by using the Chiralpak@IE column (see ESI for conditions).[#] The chiroptical properties (ECD and MR)[§] of enantioenriched **3b** samples were then studied. Figure 2 shows mirror-image spectra in CH₂Cl₂ corresponding to (M,R_{ir}) - and (P,S_{ir}) -**3b**. Complex (P,S_{ir}) -**3b**

displays the typical strong negative ECD band at 266 nm ($\Delta\epsilon = -164$) accompanied by a shoulder at 254 nm ($\Delta\epsilon = -151$), a strong positive ECD-active band at 340 nm ($\Delta\epsilon = +136$) with a shoulder at 362 nm ($\Delta\epsilon = +75$) and a weaker positive band at 304 nm ($\Delta\epsilon = +40$). Overall the ECD intensity of the chiral Ir complex **3b** is strong, as are its optical rotation values ((P,S_{ir}) -**3b**: $[\alpha]_D^{23} = +1490$, $[\phi]_D^{23} = +12350$ ($\pm 5\%$), C 1.1 10^{-4} M, CH₂Cl₂; calc. B3LYP: 13409). The question arose whether the (M,R_{ir}) - and (P,S_{ir}) -**3b** stereoisomers were the only species present in solution, *i.e.* whether the stereogenic Ir center in **3b** is configurationally stable. First of all, only one set of signals was observed in the ¹H and ¹³C NMR spectra as well as in the chiral HPLC chromatograms (see ESI). Furthermore, a sample of (M,R_{ir}) -**3b** was heated in refluxing chloroform for 2 hrs and the NMR showed no difference, suggesting that the compound is both diastereoselectively and chemically stable under these conditions (see Figure S11).[‡] Accordingly, the configuration around the Ir(III) center in **3b** is more stable than in **3a**. Note also that the (P,S_{ir}) -**3b** stereoisomer was calculated to be 1 kcal/mol lower in energy than the corresponding (P,R_{ir}) system, which may translate into a 85/15 Boltzmann ratio in favor of (P,S_{ir}) at 20°C. The experimental observations suggest strongly, however, that the formation and stability of the complexes is dictated by kinetics. Finally, the calculated chiroptical properties of (P,S_{ir}) -**3b** agree very well with experiment (Figure 3); the (P,R_{ir}) -**3b** spectrum appears similar but is somewhat more intense (ESI). The energy and assignment of the very first calculated excitation are the same for both $(P,S_{ir})/(P,R_{ir})$ -**3b** and S_{ir}/R_{ir} -**3a**. For the S_{ir} (R_{ir}) configuration it has positive (negative) rotatory strength, which may additionally support the **3a** configuration assignment. A corresponding weak band or shoulder in the broadened simulated or the experimental ECD spectrum is not visible for $(P,S_{ir})/(P,R_{ir})$ -**3b**, however, because the first excitation is over-powered by very intense higher excitations. An MO-pair analysis of the intense excitations of (P,S_{ir}) -**3b** reveals a strong involvement of the Ir(III) orbitals and carbenic π -system in the ECD-active transitions. For example, the (P,S_{ir}) -**3b** excitation with the strongest rotatory strength (no. 6, see inset of Figure 3) affords contributions from the HOMO-2-LUMO pair, which corresponds to a helicene π - π^* transition enhanced by the involvement of metal 5d orbitals, and from the HOMO-LUMO+1 pair, which has predominantly ML and ligand-to-ligand (LL) CT character. Two other excitations of moderate intensity, no. 5 and no. 1 are assigned as a mixture of ML/LL CT and LF transitions with some LM and ML CT and clearly involve the carbenic π -system. These excitations are responsible for the pronounced low-energy tail of the ECD spectrum of **3b**. These results account for the noticeable differences between the calculated spectrum of (P,S_{ir}) -**3b** versus that of the NHC-helicene ligand P -**4b**; the latter essentially affords pure π - π^* helicene character (see ESI). Overall, the spectral analysis shows that the 6-membered iridacycle affords electronic coupling between the metal center and the helicenic NHC ligand. To our knowledge this is the first time that the NHC involvement in the chiroptical

properties is identified in a chiral NHC transition metal complex.

In conclusion, we have prepared and characterized the first chiral complexes bearing a NHC-helicene ligand. These Ir(III) systems are also chiral at the metal and display efficient chiral induction from the helicene to the iridium center in the case of the NHC-[6]helicene ligand, thus giving access to complexes of well-defined stereochemistry, (*M,R_{ir}*)-(-) and (*P,S_{ir}*)-(+)-**3b**. In addition, we have shown substantial involvement of occupied MOs with electron density in the NHC group in the transitions causing the electronic circular dichroism. The use of these unprecedented Ir complexes as chiral materials and as enantioselective catalysts is under investigation. Finally, this study, combining N-heterocyclic carbenes, helicenes, and chiral systems chemistry, opens up a new field of research since many different helicene-NHC-based chiral complexes may be targeted.

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Notes and references

‡ Shoulders are present in the HPLC chromatograms and probably account for the presence of some residual isostructural (**4a/4b**)IrClCp* impurity, that is also seen by mass spectrometry (see ESI).

£ The crude NMR of pure racemic (**4b**)IrClCp*, prepared from the chloride salt of **2b**, suggests that the reaction was diastereoselective.

§ All $\Delta\epsilon$ values in this work are given in units of $M^{-1} cm^{-1}$, all specific rotations in degree $[dm (g cm^{-3})]^{-1}$, and all molar rotations in degree $cm^2 dmol^{-1}$.

It is not feasible to first generate the carbene **4b** and resolve it into its pure enantiomers, since the free **4b** is expected to be unstable.

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